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=> s dmxaa

L1 3 DMXAA

=> d 11 1-3

L1 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN

RN 853799-58-3 REGISTRY

ED Entered STN: 05 Jul 2005

CN 9H-Xanthene-4-acetic acid, 5,6-dimethyl-9-oxo-, mixt. with 2-[(2,6-dichlorophenyl)amino]benzeneacetic acid (9CI) (CA INDEX NAME) OTHER NAMES:

CN DMXAA-diclofenac mixture

MF C17 H14 O4 . C14 H11 C12 N O2

CI MXS

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPAT2, USPATFULL

CM 1

CRN 117570-53-3 CMF C17 H14 O4

CM 2

CRN 15307-86-5

CMF C14 H11 C12 N O2

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L1 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN

RN 129095-08-5 REGISTRY

ED Entered STN: 31 Aug 1990

CN 9H-Xanthene-4-acetic acid, 5,6-dimethyl-9-oxo-, sodium salt (1:1) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 9H-Xanthene-4-acetic acid, 5,6-dimethyl-9-oxo-, sodium salt (9CI)

OTHER NAMES:

CN DMXAA sodium salt

MF C17 H14 O4 . Na

CI COM

SR CA

LC STN Files: BEILSTEIN\*, CA, CAPLUS, IMSPATENTS, IMSRESEARCH, TOXCENTER, USPAT2, USPATFULL

(\*File contains numerically searchable property data)

CRN (117570-53-3)

● Na

6 REFERENCES IN FILE CA (1907 TO DATE)

6 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L1 ANSWER 3 OF 3 REGISTRY COPYRIGHT 2009 ACS on STN

RN 117570-53-3 REGISTRY

ED Entered STN: 18 Nov 1988

CN 9H-Xanthene-4-acetic acid, 5,6-dimethyl-9-oxo- (CA INDEX NAME)

OTHER NAMES:

CN 5,6-Dimethyl-9-oxo-9H-xanthen-4-ylacetic acid

CN 5,6-Dimethylxanthenone-4-acetic acid

CN AS 1404

CN ASA 404

CN DMXAA

CN NSC 640488

CN Vadimezan

MF C17 H14 O4

CI COM

SR CA

LC STN Files: ADISINSIGHT, ADISNEWS, ANABSTR, BEILSTEIN\*, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, CHEMINFORMRX, CIN, EMBASE, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, PHAR, PROMT, PROUSDDR, RTECS\*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL

(\*File contains numerically searchable property data)

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

196 REFERENCES IN FILE CA (1907 TO DATE)
4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
196 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s gemcitabine L2 11 GEMCITABINE

=> file caplus
COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 17.33 17.55

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 16:11:09 ON 19 OCT 2009 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 19 Oct 2009 VOL 151 ISS 17 FILE LAST UPDATED: 18 Oct 2009 (20091018/ED) REVISED CLASS FIELDS (/NCL) LAST RELOADED: Aug 2009 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Aug 2009

CAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2009.

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This file contains CAS Registry Numbers for easy and accurate

substance identification.

=> s 11 L3 197 L1 => s 12 5378 L2 L4 $\Rightarrow$  s 11 and 12 197 L1 5378 L2 L5 6 L1 AND L2 => dup rem 15 PROCESSING COMPLETED FOR L5 6 DUP REM L5 (0 DUPLICATES REMOVED) => d 16 1-6 ibib abs DOCUMENT NUMBER:

L6 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2009:739059 CAPLUS

151:86657

Combinations of therapeutic agents comprising vascular TITLE:

disrupting agent such as

5,6-dimethylxanthenone-4-acetic acid, for treating

cancer

Evans, Dean Brent; Jacques, Christian J. INVENTOR(S):

PATENT ASSIGNEE(S): Novartis A.-G., Switz. PCT Int. Appl., 57pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.					KIN				APPLICATION NO.						DATE			
	WO 2009076170									WO 2008-US85535					20081204				
	WO	₩:	AE, CA, FI, KG, ME, TM, AT,	AG, CH, GB, KM, MG, TT, TN, BE,	AL, CN, GD, KN, MK, RO, TR, BG,	AM, CO, GE, KP, MN, RS, TT, CH,	AO, CR, GH, KR, MW, RU, TZ, CY,	AT, CU, GM, KZ, MX, SC, UA, CZ,	AU, CZ, GT, LA, MY, SD, UG, DE,	AZ, DE, HN, LC, MZ, SE, US, DK,	DK, HR, LK, NA, SG, UZ, EE,	DM, HU, LR, NG, SK, VC, ES,	DO, ID, LS, NI, SL, VN, FI,	DZ, IL, LT, NO, SM, ZA, FR,	EC, IN, LU, NZ, ST, ZM, GB,	EE, IS, LY, OM, SV, ZW GR,	EG, JP, MA, PG, SY,	ES, KE, MD, PH, TJ,	
PRIONAB	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA  PRIORITY APPLN. INFO.:  US 2007-13335P P 20071213  AB The invention relates to a combination comprising vascular disrupting agent (VDA), such as 5,6-dimethylxanthenone-4-acetic acid or a pharmaceutically acceptable salt, ester or prodrug thereof; and one or more pharmaceutically active agents; pharmaceutical compns. comprising said combination; methods of treatment comprising said combination; processes for making said combination; and a com. package comprising said combination. Thus, the effects of 5,6-dimethylxanthenone-4-acetic acid (Compound A), trastuzumab and paclitaxel are evaluated for their antitumor activity using the BT-474 human breast ductal carcinoma xenograft model;																		
	the	e dat	a sh	ows ·	that	Com	ooun	d A	at 2	0 mg	/kg (	give	n i.	v. o:	n da	ys 1,	, 5	and 9 is	

able to produce inhibition of tumor growth; paclitaxel combined with trastuzumab is also active resulting in a combination effect; when Compound A at 20 mg/kg is combined with paclitaxel and trastuzumab, increased activity is apparent resulting in tumor regressions; using the Clark Combination Index method, synergy is indicated; the tolerability of the triple combinations is no worse than that observed when Compound A is dosed alone.

L6 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:1250046 CAPLUS

DOCUMENT NUMBER: 149:448110

TITLE: Preparation of Iso CA-4 and analogs as potent

cytotoxic agents and inhibitors of polymerization of

tubulin

INVENTOR(S): Alami, Mouad; Brion, Jean-Daniel; Provot, Olivier;

Peyrat, Jean-Francois; Messaoudi, Samir; Hamze, Abdallah; Giraud, Anne; Bignon, Jerome; Bakala,

Joanna; Liu, Jian-Miao

PATENT ASSIGNEE(S): Centre National De La Recherche Scientifique, Fr.

SOURCE: PCT Int. Appl., 78pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: French

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

GΙ

PATENT	PATENT NO.					DATE APPLICATION NO.					DATE						
WO 200	WO 2008122620			A1 2008			081016 WO 2008-EP5					118 20080404				404	
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	FΙ,	GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	
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RW	: AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FΙ,	FR,	GB,	GR,	HR,	HU,	
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	ΑM,	ΑZ,	BY,	KG,	KΖ,	MD,	RU,	ТJ,	$_{ m MT}$								
FR 291	FR 2914640					A1 20081010			FR 2007-54280					20070404			
PRIORITY AP	PRIORITY APPLN. INFO.:								FR 2	007-	5428	0	i	A 2	0070	404	
OTHER SOURC	MAR:	PAT	149:	4481	10												

AB Isocombretastatin A-4 and analogs I [R1, R2, R3 = methoxy (possibly substituted by one or more fluorine atoms); R5 = R6 = hydrogen or fluorine; A = ring chosen from (un)substituted aryls and heteroaryls].

The process for the preparation of I comprises: (a) reaction of acetophenone derivative II with an organometallic compound, A-M [M = alkali metal or earth alkaline metal substituted with a halogen]; and (b) reaction of the resulting phenylethanol derivative III with an acid to form I. Thus, Iso-CA-4 [I; A = C6H3OH-3-OMe-4, R1 = R2 = R3 = OMe, R4 = R5 = R6 = H (IV)] was prepared from 3,4,5-trimethoxyacetophenone (II; R1 = R2 = R3 = OMe, R4 = R5 = R6 = H) via reaction in PhMe with tert-butyl(5-lithio-2-methoxyphenoxy)dimethylsilane [prepared from tert-butyl(5-iodo-2-methoxyphenoxy)dimethylsilane via lithiation with Me3CLi in hexane], dehydration of III with p-toluenesulfonic acid in CH2Cl2, and desilylation with K2CO3 in MeOH. The cytotoxic activity of IV was determined [IC50 = 2-4 nM vs. HCT116; IC50 = 5 nM vs. K562 cells; IC50 = 2 nM vs. B16F10 cells; IC50 = 8 nM vs. U87 cells; IC50 = 8 nM vs. A549 cells; IC50 = 4.5 nM vs. M435 cells; IC50 = 4 nM vs. M231 cells; IC50 = 2.2  $\mu$ M vs tubulin polymerization].

OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:473431 CAPLUS

DOCUMENT NUMBER: 148:463206

TITLE: oncolytic viruses and antiangiogenic agents in the

treatment of cancer

INVENTOR(S): Karrasch, Matthias; Mescheder, Axel

PATENT ASSIGNEE(S): Medigene AG, Germany SOURCE: PCT Int. Appl., 69pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

P	PATENT NO. 			KIND DATE				•	APPL	ICAT		DATE					
W(				A1 20080417			0417	WO 2007-EP8930					20071015				
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		KM,	KN,	KP,	KR,	KΖ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ME,
		MG,	MK,	MN,	MW,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,
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		TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW				
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El	EP 2073823				A1		2009	0701	EP 2007-819001					20071015			
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		AL,	BA,	HR,	MK,	RS											
PRIORI:	RIORITY APPLN. INFO.:									US 2 WO 2					P 2	0061 0071	

AB The invention relates to a combination of at least one oncolytic virus and at least one antiangiogenic agent and to the use of this combination in tumor therapy. Intraarterial infusions of oncolytic virus NV1020 to a patient with progressive metastatic colorectal adenocarcinoma followed by CPT-11 plus cetuximab resulted in stabilization of the disease at 6 mo post treatment.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS

L6 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:984120 CAPLUS

DOCUMENT NUMBER: 143:279360

TITLE: Methods of detecting CD133 antigen (AC133) expression

level and use as biomarker for human cancer diagnosis

and therapy monitor

INVENTOR(S): Penning, Maarten Tjerk; Van den Broek, Sebastiaan

Johannes Jacobus; Voest, Emile Eugene; Beerepoot,

Laurens Victor; Mehra, Niven

PATENT ASSIGNEE(S): Primagen Holding B. V., Neth.; UMC Utrecht Holding B.

V.

SOURCE: PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

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PATENT NO.
                            KIND DATE
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                                    20050909 WO 2005-NL155
                                                                               20050302
     WO 2005083123
                             A1
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PRIORITY APPLN. INFO.:
                                                    EP 2004-75686
                                                                          A 20040302
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                                                    US 2004-549450P
                                                    EP 2005-710924
                                                                          A 20050302
                                                                           W 20050302
                                                    WO 2005-NL155
                                                    US 2006-514345 B1 20060831
     This invention provides methods of detecting CD133 antigen (AC133)
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AB This invention provides methods of detecting CD133 antigen (AC133) expression level and use as a biomarker for human cancer diagnosis and therapy monitor. Blood anal. including number of circulating endothelial cells and expression levels of human genes AC133 (CD133), EST032 and U1A evaluated by NASBA anal., were determined prior to and during chemotherapy using drugs such as angiostatin or PrimMed01, gemcitabine, and cisplatin, for a wide range of human tumor types. A use of a nucleic acid mol. comprising at least part of a sequence of AC133 or an analog thereof for monitoring a treatment of an individual suffering from a disease is also provided, as well as a diagnostic kit comprising such nucleic acid mol.

REFERENCE COUNT:

9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:975665 CAPLUS

DOCUMENT NUMBER: 143:264929

TITLE: Methods for detecting AC133 antigen mRNA for diagnosis

and treatment of cancer and other diseases

INVENTOR(S): Penning, Maarten Tjerk; Beerepoot, Laurens Victor; Van

Den Broek, Sebastiaan Johannes Jacobus; Mehra, Niven;

Voest, Emile Eugene

PATENT ASSIGNEE(S): Primagen Holding B.V., Neth.; UMC Utrecht Holding B.V.

SOURCE: Eur. Pat. Appl., 28 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

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PATENT NO.
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PRIORITY APPLN. INFO.:
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AB
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AB The invention provides methods for detecting AC133 antigen mRNA for diagnosis and treatment of cancer and other diseases. AC133 antigen mRNA may be quantitated by PCR, RT-PCR, NASBA, SDA, TMA, bDNA or rolling circle amplification. Diseases include cancer and heart disease, high blood pressure, ischemia, stroke, psoriasis, Crohn's disease, rheumatoid arthritis, endometriosis, atherosclerosis, obesity, diabetes mellitus, diabetic retinopathy, macular degeneration, Alzheimer's disease, Peutz Jegher's syndrome, multiple sclerosis, systemic lupus erythematosus, Wegener's granulomatosis, vasculitis, sickle cell disease, thalassemia and angina.

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ACCESSION NUMBER: 2003:202462 CAPLUS

DOCUMENT NUMBER: 138:226761

TITLE: Synergistic anticancer combinations containing

5,6-dimethylxanthenone-4-acetic acid

Wilson, William Robert; Siim, Bronwyn Gae Cancer Research Technology Limited, UK

PATENT ASSIGNEE(S): Cancer Research Technology Li

PCT Int. Appl., 31 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

INVENTOR(S):

SOURCE:

PATEN	NT NO.	KIND	DATE	APPLICATION NO.				
WO 20	003020259 003020259	A2		WO 2002-GB4025				
V	CO, CR, C GM, HR, H LS, LT, L	CU, CZ, I IU, ID, I LU, LV, N	DE, DK, DM, IL, IN, IS, MA, MD, MG,	BA, BB, BG, BR, BY, DZ, EC, EE, ES, FI, JP, KE, KG, KP, KR, MK, MN, MW, MX, MZ, SI, SK, SL, TJ, TM,	GB, GD, GE, GH, KZ, LC, LK, LR, NO, NZ, OM, PH,			
F	UA, UG, U RW: GH, GM, K KG, KZ, M FI, FR, G	JS, UZ, V CE, LS, M ID, RU, T GB, GR, J	VC, VN, YU, MW, MZ, SD, IJ, TM, AT, IE, IT, LU,		ZW, AM, AZ, BY, DE, DK, EE, ES, TR, BF, BJ, CF,			
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EP 14	423105	B1	20040602 20081203 DK, ES, FR,	EP 2002-758562  GB, GR, IT, LI, LU,				
JP 20 CN 17 NZ 53 EP 17	IE, SI, I 002012258 005509599 708296 31045	T, LV, E A T A A A	FI, RO, MK, 20041019 20050414	CY, AL, TR, BG, CZ, BR 2002-12258 JP 2003-524567 CN 2002-817257 NZ 2002-531045	EE, SK  20020903 20020903 20020903 20020903			
NZ 54 CN 19 NZ 55 AT 41 ES 23 NO 20 ZA 20 US 20 MX 20 IN 20 AU 20 US 20	LI, LU, M 46573 994287 54093 15963 321283 004000591 004001078 0040204480 004002004	IC, NL, E A A A T T3 A A A1 A A1	PT, SE, SK, 20070531 20070711 20080731 20081215 20090604 20040430 20050415 20041014	MX 2004-2004	RO, SI  20020903 20020903 20020903 20020903 20040210 20040210 20040302 20040302 20040402 20070509			

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The present invention relates to synergistic combinations of the AB 5,6-dimethylxanthenone-4-acetic acid (DMXAA) and a compound selected from platinum compds., Vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors, which have antitumor activity. More particularly, the invention is concerned with the use of such combinations in the treatment of cancer and pharmaceutical compds. containing the combinations. The antitumor activity and host toxicity of DMXAA/cytotoxic drug combinations was assessed by varying the dose of chemotherapeutic drug up to the toxicity limit, with co-administration of a fixed DMXAA dose (80  $\mu$ mol/kg, ca. 80% of MTD), and evaluating subsequent tumor growth delay. Of the 7 drugs investigated, 4 (doxorubicin, 5-fluorouracil, cyclophosphamide and cisplatin) had appreciable activity against this tumor as indicated by dose-response relationships providing significant slopes by linear regression, and highly significant growth delays of 10 days at their MTDs.

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